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	Application Number	10/564,614	
INFORMATION DISCLOSURE	Filing Date	May 30, 2006	
STATEMENT BY APPLICANT	First Named Inventor	Vladimir Subr	
(Use as many sheets as necessary)	Art Unit	4161	
(Use as many sneets as necessary)	Examiner Name	Kevin S. Orwig	
Sheet 1 of 2	Attorney Docket Number	J126-022 US	

		NON PATENT LITERATURE DOCUMENTS	
Examination Initials*	er Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T
		K. Ulbrich, et al. "Polymeric drugs based on conjugates of synthetic and natural macromolecules I. Synthesis and physico-chemical characterisation" J. Controlled Release 64, 2000, 63-79	
***************************************	***************************************	B. Rihova, et al. " Polymeric drugs based on conjugates of synthetic and natural macromolecules II. Anticancer activity of antibody or (Eab') sub 3 targeted conjugates and combined therapy with	
		immunomodulators" J. Controlled Release 64 (2000) 241-261	<u></u>
	000000000000000000000000000000000000000	J. Kopecek, et al. " HPMA copolymer-anticancer drug conjugates: design, activity, and mechanism	ļ
		of action"	
		K. Ulbrich, et al. "Polymeric Conjugates of Drugs and Antibodies for Site-Specific	
000000000000000000000000000000000000000	000000000000000000000000000000000000000	Drug Delivery, Macromol. Symp*	0000000
		J. Kopecek, et al., "HPMA copolymer-anticancer drug conjugates: design, activity, and	
		mechanism of action"	
000000000000000000000000000000000000000	***************************************	R.A. Vasay, et al., "HPMA co-polymer doxorubicio".	
			L
		P.A. Vasey, et al., "Phase I clinical and pharmacokinetic study of PK1 (W-(2-hydroxypropyl)methacrylamide copolymer doxorubicin). First member of a new class of	<u> </u>
		chemotherapeutic	
		P. J. Julyan, et al., "Preliminary clinical study of the distribution of HPMA copolymers bearing	
	000000000000000000000000000000000000000	doxorubicin and galaciosamine , J. Comrolled Release 57 (1999) 281-290	
		A. H. Thomson et al., "Population pharmacokinetics in phase I drug development; a phase I study	
		of PKI in patients with solid tumors"	
		L. W. Seymour et al., Hepatic drug targeting: Phase Levaluation of polymer bound	
		doxorubicin.	

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Examiner	Wayin Onvial		Date	03/08/2009
Signature	/Kevin Orwig/	·	Considered	03/00/2009

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			J. Clin, "Phase I clinical and pharmacokinetic study of PNU 166945, a novel water-soluble	
000000000000000000000000000000000000000			prodrug of paclitaxel"	
000000000000000000000000000000000000000		00000000000000	M. I.M. Tenwogt, "Phase I clinical and pharmacokinetic study of PNI L166945, a novel water-soluble	
			polymer-conjugated prodrug of paclitaxel. Anti-Cancer Drugs	
			M. Bouma, Stability and compatiblity of the investigational polymer-conjugated platinum anticancer	
200000000000000000000000000000000000000			agent AP 5280 in intusion systems and its nemolytic potential	
			M.M. Tibben, "Determination of total platinum in plasma and plasma ultrafiltrate, from subjects	
100000000000000000000000000000000000000		000000000000000000000000000000000000000	dosed with the platinum-containing N-(2-hydroxypropyr)methacrylamide copolymer AP5200; by use of graphite-furnace Zeeman atomic-absorption"	50000000000000
***************************************	***************************************	***************************************	B. Rihova et al. "Drug-HPMA-Hulg conjugates effective against human solid cancer"	
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